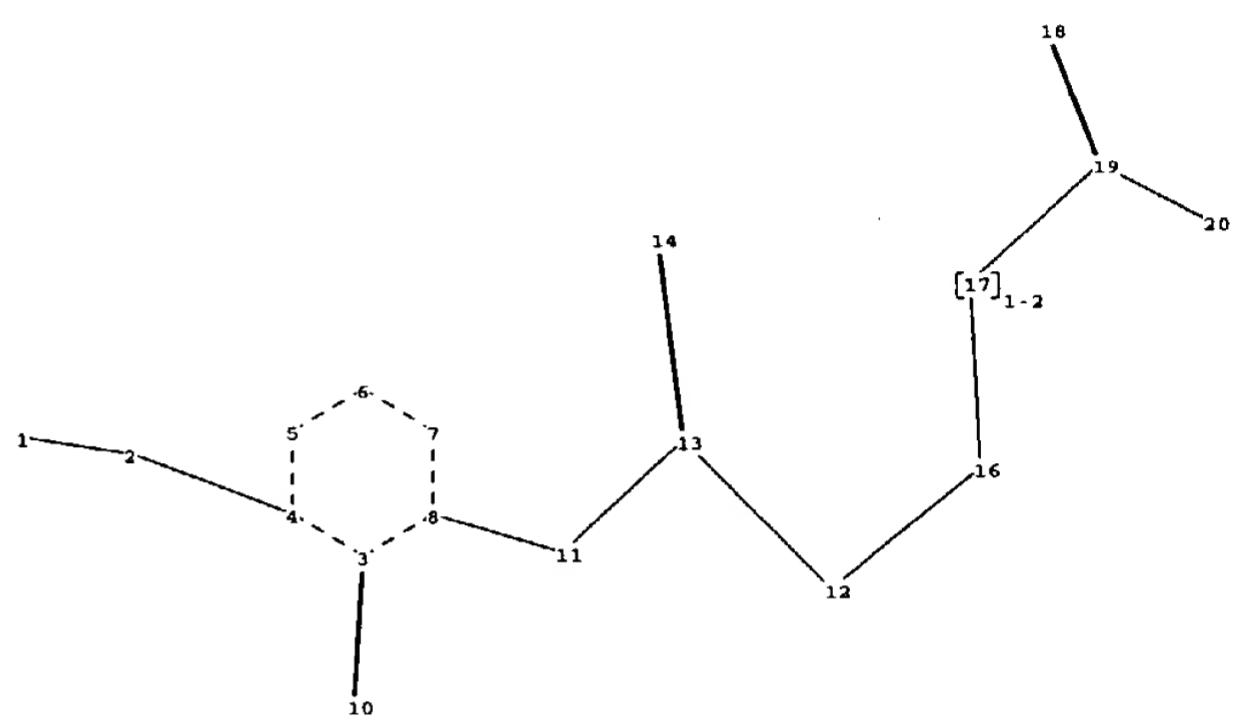
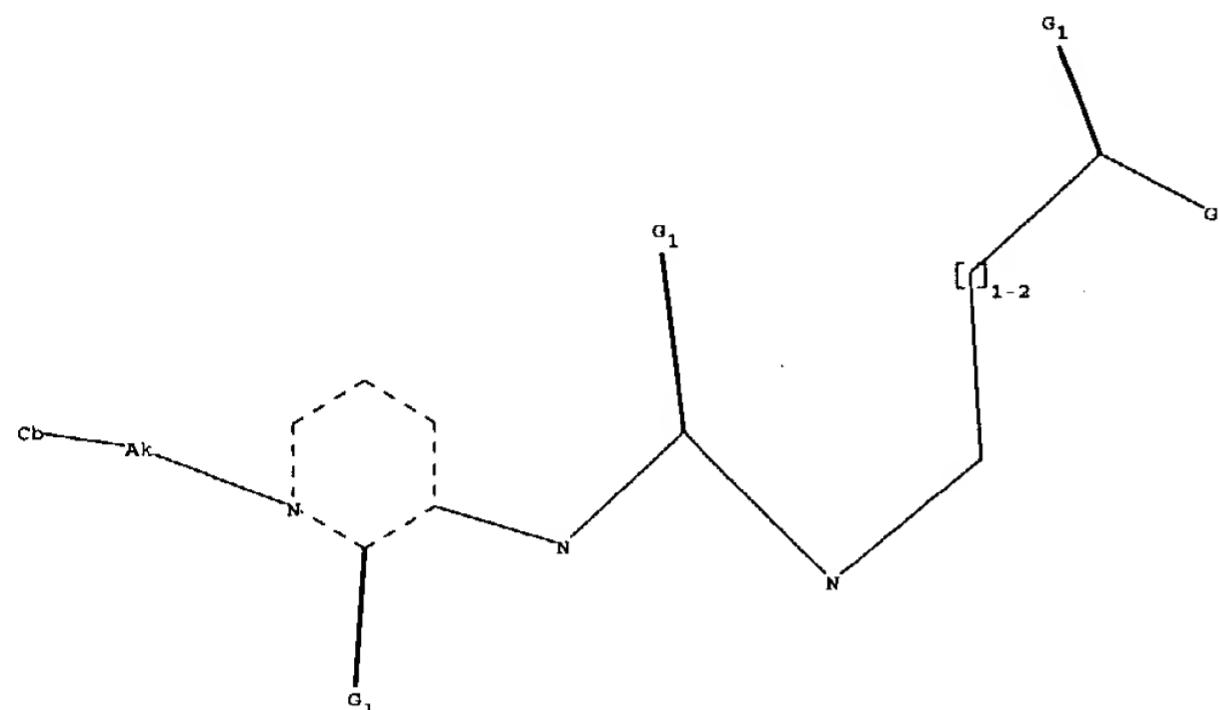


38 52
75

chain nodes :

1 2 10 11 12 13 14 16 17 18 19 20

ring nodes :

3 4 5 6 7 8

chain bonds :

1-2 2-4 3-10 8-11 11-13 12-13 12-16 13-14 16-17 17-19 18-19 19-20

ring bonds :

3-4 3-8 4-5 5-6 6-7 7-8

exact/norm bonds :

1-2 2-4 3-4 3-8 3-10 4-5 5-6 6-7 7-8 8-11 11-13 12-13 12-16 13-14 18-19
19-20

exact bonds :

16-17 17-19

isolated ring systems :

containing 3 :

G1:0,S

Match level :

1:Atom 2:CLASS 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 10:CLASS 11:CLASS
12:CLASS 13:CLASS 14:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS 20:CLASS

Patent Assignment Abstract of Title

Total Assignments: 1

Application #: 09973142 **Filing Dt:** 10/09/2001

Patent #: NONE

Issue Dt:

PCT #: NONE

Publication #: 20040063955

Pub Dt: 04/01/200

Inventors: Ronald J. Biediger, Qi Chen, E. Radford Decker, George W. Holland, Jamal M. Kassir, Wen Li, Robert V. Market, Ian L. Scott, Chengde Wu, Jian Li

Title: Carboxylic acid derivatives that inhibit the binding of integrins to their receptors

Assignment: 1

Reel/Frame: <u>012249/0414</u>	Received: 10/18/2001	Recorded: 10/09/2001	Mailed: 12/20/2001	Pages: 2
---------------------------------------	--------------------------------	--------------------------------	------------------------------	-----------------

Conveyance: ASSIGNMENT OF ASSIGNORS INTEREST (SEE DOCUMENT FOR DETAILS).

Assignors: <u>BIEDIGER, RONALD J.</u>	Exec Dt: 10/02/2001
<u>CHEN, QI</u>	Exec Dt: 10/02/2001
<u>DECKER, RADFORD E.</u>	Exec Dt: 10/02/2001
<u>HOLLAND, GEORGE W.</u>	Exec Dt: 10/02/2001
<u>KASSIR, JAMAL M.</u>	Exec Dt: 10/02/2001
<u>LI, WEN</u>	Exec Dt: 10/02/2001
<u>MARKET, ROBERT V.</u>	Exec Dt: 10/02/2001
<u>SCOTT, IAN L.</u>	Exec Dt: 10/04/2001
<u>WU, CHENGDE</u>	Exec Dt: 10/02/2001
<u>LI, JIAN</u>	Exec Dt: 10/02/2001

Assignee: TEXAS BIOTECHNOLOGY CORPORATION

7000 FANNIN, SUITE 1920
HOUSTON, TEXAS 77030

Correspondent: ROCKY, MILNAMOW & KATZ, LTD.

MARTIN L. KATZ
TWO PRUDENTIAL PLAZA
180 NORTH STETSON AVENUE, SUITE 4700
CHICAGO, IL 60601

Search Results as of: 4/11/2004 5:41:04 P.M.

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Web interface last modified: Oct. 5, 2002

<u>NEWS 1</u>	Web Page URLs for STN Seminar Schedule - N. America
<u>NEWS 2</u>	"Ask CAS" for self-help around the clock
<u>NEWS 3</u>	NOV 24 MSDS-CCOHS file reloaded
<u>NEWS 4</u>	DEC 08 CABA reloaded with left truncation
<u>NEWS 5</u>	DEC 08 IMS file names changed
<u>NEWS 6</u>	DEC 17 DGENE: Two new display fields added
<u>NEWS 7</u>	DEC 18 BIOTECHNO no longer updated
<u>NEWS 8</u>	DEC 19 CROPU no longer updated; subscriber discount no longer available
<u>NEWS 9</u>	DEC 22 ABI-INFORM now available on STN
<u>NEWS 10</u>	JAN 27 Source of Registration (SR) information in REGISTRY updated and searchable
<u>NEWS 11</u>	JAN 27 A new search aid, the Company Name Thesaurus, available in CA/CAplus
<u>NEWS 12</u>	FEB 05 German (DE) application and patent publication number format changes
<u>NEWS 13</u>	MAR 03 MEDLINE and LMEDLINE reloaded
<u>NEWS 14</u>	MAR 03 MEDLINE file segment of TOXCENTER reloaded
<u>NEWS 15</u>	MAR 03 FRANCEPAT now available on STN
<u>NEWS 16</u>	MAR 29 Pharmaceutical Substances (PS) now available on STN
<u>NEWS 17</u>	MAR 29 WPIFV now available on STN
<u>NEWS 18</u>	MAR 29 No connect hour charges in WPIFV until May 1, 2004
<u>NEWS 19</u>	MAR 29 New monthly current-awareness alert (SDI) frequency in RAPRA
<u>NEWS EXPRESS</u>	MARCH 31 CURRENT WINDOWS VERSION IS V7.00A, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 3 MARCH 2004
<u>NEWS HOURS</u>	STN Operating Hours Plus Help Desk Availability
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<u>NEWS WWW</u>	CAS World Wide Web Site (general information)

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=> file reg
COST IN U.S. DOLLARS
SINCE FILE
ENTRY
TOTAL
SESSION
0.21
0.21
FULL ESTIMATED COST

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STRUCTURE FILE UPDATES: 9 APR 2004 HIGHEST RN 673855-15-7
 DICTIONARY FILE UPDATES: 9 APR 2004 HIGHEST RN 673855-15-7

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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=>
 L1 STRUCTURE UPLOADED

=> d 11
 L1 HAS NO ANSWERS
 L1 STR

=> s 11
 SAMPLE SEARCH INITIATED 17:38:06 FILE 'REGISTRY'
 SAMPLE SCREEN SEARCH COMPLETED - 45 TO ITERATE

100.0% PROCESSED 45 ITERATIONS 14 ANSWERS
 SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
 BATCH **COMPLETE**
 PROJECTED ITERATIONS: 498 TO 1302
 PROJECTED ANSWERS: 56 TO 504

L2 14 SEA SSS SAM L1

=> s 11 full
 THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 155.00 U.S. DOLLARS
 DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y
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 FULL SCREEN SEARCH COMPLETED - 1111 TO ITERATE

100.0% PROCESSED 1111 ITERATIONS 440 ANSWERS
 SEARCH TIME: 00.00.01

L3 440 SEA SSS FUL L1

=> file hcaplus
 COST IN U.S. DOLLARS SINCE FILE TOTAL
 ENTRY SESSION
 FULL ESTIMATED COST 157.10 157.31

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FILE COVERS 1907 - 11 Apr 2004 VOL 140 ISS 16
FILE LAST UPDATED: 9 Apr 2004 (20040409/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13
L4 2 L3

=> d 14, ibib abs fhitstr, 1-2

L4 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2004 ACS on STN

Full Text	Citing References
ACCESSION NUMBER: 2002:349146 HCAPLUS	
DOCUMENT NUMBER: 136:369608	
TITLE: Preparation of 3-(N'-oxodihydropyridinylureido)-3-phenylpropanoates as inhibitors of $\alpha 4\beta 1$ integrin binding	
INVENTOR(S): Biediger, Ronald J.; Chen, Qi; Holland, George W.; Kassir, Jamal M.; Li, Wen; Market, Robert V.; Scott, Ian L.; Wu, Chengde; Decker, Radford E.; Li, Jian	
PATENT ASSIGNEE(S): Texas Biotechnology Corporation, USA	
SOURCE: Eur. Pat. Appl., 131 pp.	
DOCUMENT TYPE: Patent	
LANGUAGE: English	
FAMILY ACC. NUM. COUNT: 3	
<u>PATENT INFORMATION:</u>	

No Same
app-

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1203766	A2	20020508	EP 2001-125494	20011106
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
US 2004063955	A1	20040401	US 2001-973142	20011009
ZA 2001008777	A	20030124	ZA 2001-8777	20011024
<u>PRIORITY APPLN. INFO.:</u>				
US 2000-707068 A 20001106				
US 2001-973142 A 20011009				
US 1999-132971P P 19990507				
US 2000-565920 A2 20000505				

OTHER SOURCE(S): MARPAT 136:369608
AB Title compds. were prep'd. Thus, 2-ClC₆H₄CH₂ZNH₂ (Z = 4-ethyl-2-oxo-1,2-dihydropyridine-1,3-diyl) (prepn. given) was condensed with (S)-4-MeC₆H₄CH(NH₂)CH₂CO₂Et and COCl₂ to give, after sapon., (S)-2-ClC₆H₄CH₂ZNHCONHCH(C₆H₄Me-4)CH₂CO₂H (Z as above). Data for biol. activity of title compds. were given.

IT 307520-20-3P

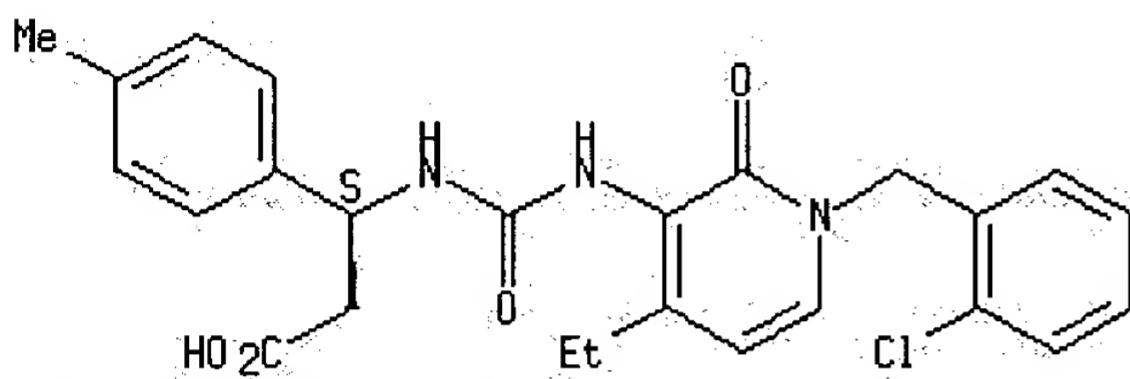
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of 3-(N'-oxodihydropyridinylureido)-3-phenylpropanoates as inhibitors of $\alpha 4\beta 1$ integrin binding)

RN 307520-20-3 HCAPLUS

CN Benzenepropanoic acid, β -[[[[1-[(2-chlorophenyl)methyl]-4-ethyl-1,2-dihydro-2-oxo-3-pyridinyl]amino]carbonyl]amino]-4-methyl-, (β S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L4 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2004 ACS on STN

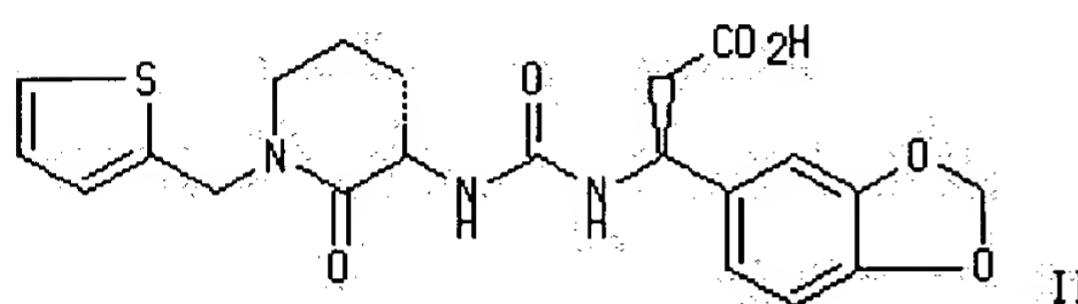
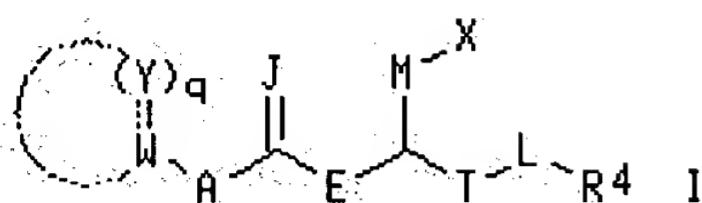
Full Text Citing References

ACCESSION NUMBER: 2000:814302 HCAPLUS
 DOCUMENT NUMBER: 133:362963
 TITLE: Preparation of β -amino acid derivatives that inhibit the binding of integrins to their receptors
 INVENTOR(S): Biediger, Ronald J.; Chen, Qi; Holland, George W.; Kassir, Jamal M.; Li, Wen; Market, Robert V.; Scott, Ian L.; Wu, Chengde
 PATENT ASSIGNEE(S): Texas Biotechnology Corporation, USA
 SOURCE: PCT Int. Appl., 113 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

*co-pending
copending
PON?*

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000067746	A1	20001116	WO 2000-US12303	20000505
WO 2000067746	C2	20020829		
	W:	AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM		
	RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		
EP 1176956	A1	20020206	EP 2000-937527	20000505
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO		
SI 20744	C	20020630	SI 2000-20021	20000505
BR 2000010293	A	20020716	BR 2000-10293	20000505
JP 2002544161	T2	20021224	JP 2000-616772	20000505
ZA 2001008774	A	20030124	ZA 2001-8774	20011024

NO 2001005418 A 20011221 NO 2001-5418 20011106
 PRIORITY APPLN. INFO.: US 1999-132971P P 19990507
 OTHER SOURCE(S): MARPAT 133:362963 W 20000505
 GI



AB Title compds. I [Y, at each occurrence, independently = CO, N, CR1, CR2R3, NR5, CH, O, or S; q = 3-10; A = O, S, CR16R17, NR6; E = CH2, O, S, NR7; J = O, S, NR8; M = CR9R10 or (CH2)0-3; T = CO or (CH2)0-3; L = O, NR11, S, (CH2)0-1; X = CO2B, PO3H2, SO3H, SO2NH2, SO2NHCOR12, OPO3H2, CONHCOR13, CONHSO2R14, tetrazolyl, hydroxyl, H; W = C, CR15, N; B, R1-17 = H, halo, hydroxyl, alkyl, alkoxy, aliph. acyl, CF3, nitro, cycloalkyl, alkylheteroaryl, sulfonyl, carboxyl, etc.] or their pharmaceutically acceptable salts were prep'd. for inhibition of the binding of $\alpha 4\beta 1$ integrin to its receptors. Thus, II was prep'd. and assayed ($IC_{50} = 0.2 \mu M$) for its ability to suppress binding using a 26-amino acid peptide contg. the CS-1 sequence of fibronectin with N-terminal cysteine coupled to maleimide activated ovalbumin.

IT 307520-20-3P

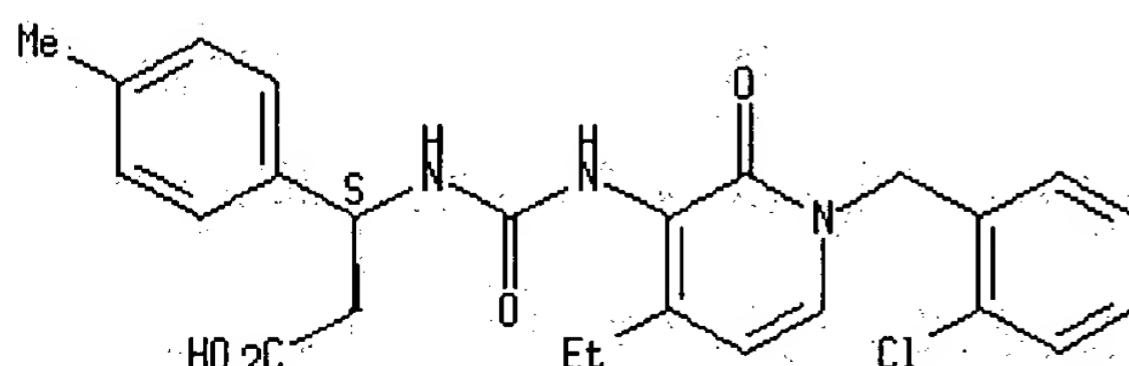
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of β -amino acid derivs. that inhibit the binding of integrins to their receptors)

RN 307520-20-3 HCPLUS

CN Benzenepropanoic acid, β -[[[[1-[(2-chlorophenyl)methyl]-4-ethyl-1,2-dihydro-2-oxo-3-pyridinyl]amino]carbonyl]amino]-4-methyl-, (β S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT:

3

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his

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FILE 'REGISTRY' ENTERED AT 17:34:59 ON 11 APR 2004

L1 STRUCTURE UPLOADED
L2 14 S L1
L3 440 S L1 FULL

FILE 'HCAPLUS' ENTERED AT 17:38:12 ON 11 APR 2004

L4 2 S L3

=> file caold

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	23.67	180.98
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-1.39	-1.39

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FILE COVERS 1907-1966

FILE LAST UPDATED: 01 May 1997 (19970501/UP)

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This file supports REGISTRY for direct browsing and searching of all substance data from the REGISTRY file. Enter HELP FIRST for more information.

=> s 13

L5 0 L3

=> file file hcaplus

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ENTER A FILE NAME OR (IGNORE):s 119:8683/dn

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SAFETY - Occupational Health and Safety Cluster

SESSION - Current files with L-numbers Cluster

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SUPPLIERS - Product Directories and Suppliers Cluster

SCISEARCH - ISI Science Citation Index from 1974 - present

SIGLE - Grey Literature in Europe from 1976 - present

SOLIDSTATE - Solid State and Superconductivity Abstracts from 1981

SOLIS - German literature in social sciences 1945-present

SPECINFO - Spectral Database Information System
 STANDARDS - The International Standards Database
 STNGUIDE - Descriptive information about STN databases
 STNMAIL - STN Electronic Mail Service
 SWETSCAN - Swets Table of Contents from 1993 - present
 SYNTHLINE - Synthline Drug Synthesis Database 1984-present
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=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.42	181.40
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CA SUBSCRIBER PRICE	0.00	-1.39

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FILE COVERS 1907 - 11 Apr 2004 VOL 140 ISS 16
 FILE LAST UPDATED: 9 Apr 2004 (20040409/ED)

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=> s 119:8683/dn
 L6 1 119:8683/DN

=> d 16, ibib abs fhitstr, 1

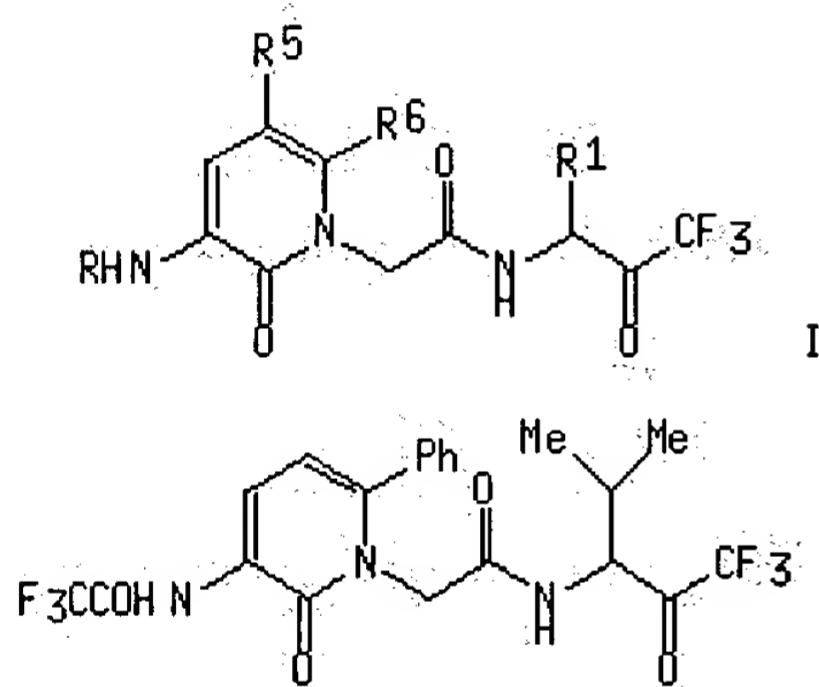
L6 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2004 ACS on STN

	Full Text	Citing References
ACCESSION NUMBER:	1993:408683 HCAPLUS	
DOCUMENT NUMBER:	119:8683	
TITLE:	Preparation of oxopyridylacetamides as human leukocyte elastase inhibitors	
INVENTOR(S):	Bernstein, Peter Robert; Shaw, Andrew; Thomas, Royston Martin; Wolanin, Donald John; Warner, Peter	
PATENT ASSIGNEE(S):	Imperial Chemical Industries PLC, UK	
SOURCE:	Eur. Pat. Appl., 96 pp.	
DOCUMENT TYPE:	Patent	
LANGUAGE:	English	
FAMILY ACC. NUM. COUNT:	3	

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 509769	A2	19921021	EP 1992-303358	19920415
EP 509769	A3	19930901		
EP 509769	B1	19960911		
NO 9201451	A	19921019	NO 1992-1451	19920410
CA 2065794	AA	19921019	CA 1992-2065794	19920410
AU 9214827	A1	19921022	AU 1992-14827	19920410
AU 660664	B2	19950706		
HU 66541	A2	19941228	HU 1992-1225	19920410
JP 06056785	A2	19940301	JP 1992-143140	19920418
<u>PRIORITY APPLN. INFO.:</u>				
			GB 1991-8357	19910418
			GB 1991-8358	19910418
			GB 1992-5392	19920312

OTHER SOURCE(S) : MARPAT 119:8683
GI



AB Title compds. [I; R = H, CHO, F₃CCO, acyl; R₁ = C₁-5 alkyl; R₅, R₆ = H, alkyl; or one of R₅, R₆ = H, Me; the other = BY; B = (substituted) (hetero)aryl; Y = bond, CH₂, CH₂CH₂, trans-CH:CH], were prep'd. Thus, title compd. II, prep'd. by oxidn. of the corresponding hydroxyamide, inhibited human leukocyte elastase with K_i = 39 nM.

=> sel rn
E1 THROUGH E629 ASSIGNED

=> s 125:599235/dn
L7 0 125:599235/DN

=> s 125:599235/dn
L8 0 125:599235/DN

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	12.06	193.46
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION

CA SUBSCRIBER PRICE	-0.69	-2.08
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FILE LAST UPDATED: 9 Apr 2004 (20040409/ED)

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=> s 125:599235/dn
L9 0 125:599235/DN

=> s 125:599235/an
L10 0 125:599235/AN

=>